

5 PHARMACEUTICAL COMPOSITIONS AND METHODS FOR THE PREVENTION
OR TREATMENT OF CARDIOVASCULAR, CARDIOPULMONARY,
PULMONARY OR RENAL DISEASES

Related Applications

10 Benefit of DE 10301371.7, filed January 16, 2003; DE 10335027.6, filed July 31, 2003; U.S. Provisional Application No. 60/446,695, filed February 11, 2003; and U.S. Provisional Application No. 60/503,317, filed September 16, 2003, are hereby claimed.

15 Background of the Invention

The invention relates to: a process for the prevention or treatment of cardiovascular, cardiopulmonary, pulmonary or renal diseases, particularly in people in whom diabetes has been diagnosed or who are suspected of prediabetes, for preventing diabetes and prediabetes, or for the treatment of
20 Metabolic Syndrome and insulin resistance in patients with normal blood pressure. The process comprises generally administering effective amounts of the angiotensin II receptor antagonist telmisartan and the HMG-CoA-reductase inhibitor atorvastatin or polymorphs or salts thereof to a person in need of treatment. The invention further relates to suitable pharmaceutical compositions
25 which contain telmisartan and atorvastatin or polymorphs or salts thereof, as a combined preparation for simultaneous, separate or sequential use in the prevention or treatment of these diseases, as well as the combined use of telmisartan and atorvastatin or polymorphs or salts thereof for preparing a pharmaceutical composition for the prevention or treatment of these diseases.

30 Angiotensin II (ANG II) plays an important part in pathophysiology, particularly as the most potent agent for increasing blood pressure in humans. It is known that in addition to its effect of raising blood pressure ANG II also has growth-promoting effects which contribute to left ventricular hypertrophy, vascular thickening,
35 atherosclerosis, kidney failure and stroke. On the other hand, bradykinin has

- 5 vasodilating and tissue-protecting effects. Therefore, ANG II antagonists are suitable for the treatment of raised blood pressure and congestive heart failure in mammals. Examples of ANG II antagonists are described in EP-A-0 502 314, EP-A-0 253 310, EP-A-0 323 841, EP-A-0 324 377, US-A-4 355 040 and US-A-4 880 804. Examples of ANG II antagonists are candesartan, eprosartan, irbesartan,
10 losartan, olmesartan, tasosartan, valsartan or telmisartan.

The antihypertensive and kidney-protecting effects of ANG II antagonists are described for example in the following publications:

- W. Wienen et al.: Antihypertensive and renoprotective effects of telmisartan
15 after long term treatment in hypertensive diabetic (D) rats, 2nd Int. Symposium on Angiotensin II Antagonism, February 15-18, 1999, The Queen Elizabeth II Conference Center, London, UK, Book of Abstracts, Abstract No. 50;
 - J. Wagner et al.: Effects of AT1 receptor blockade on blood pressure and the renin angiotensin system in spontaneously hypertensive rats of the stroke prone
20 strain, Clin. Exp. Hypertens., vol. 20 (1998), p. 205-221; and
 - M. Böhm et al.: Angiotensin II receptor blockade in TGR(mREN2)27: Effects of renin-angiotensin-system gene expression and cardiovascular functions, J. Hypertens., vol. 13 (8) (1995), p. 891-899.
- 25 Other renoprotective effects of ANG II antagonists which were found in first clinical trials are described in the following publications, for example:
- S. Andersen et al.: Renoprotective effects of angiotensin II receptor blockade in type 1 diabetic patients with diabetic nephropathy, Kidney Int., vol. 57 (2) (2000), p. 601-606;
 - 30 • L.M. Ruilope: Renoprotection and renin-angiotensin system blockade in diabetes mellitus, Am. J. Hypertens., vol. 10(12 PT 2) Suppl. (1997), p. 325-331; and
 - J.F.E. Mann: Valsartan and the kidney: Present and future, J. Cardiovasc. Pharmacol., vol. 33, Suppl. 1 (1999), p. 37-40.

- 5 Moreover the effects of ANG II antagonists on endothelial dysfunction are described in the following publications, for example:
- E.L. Schiffrin et al.: Correction of arterial structure and endothelial dysfunction in human essential hypertension by the angiotensin receptor antagonist losartan, *Circulation*, vol. 101(14) (2000), p. 1653-1659;
 - 10 • R.M. Touyz et al.: Angiotensin II stimulates DNA and protein synthesis in vascular smooth muscle cells from human arteries: role of extracellular signal-regulated kinases, *J. Hypertens.*, vol. 17(7) (1999), p. 907-916;
 - E.L. Schiffrin: Vascular remodelling and endothelial function in hypertensive patients: Effects of antihypertensive therapy, *Scand. Cardiovasc. J.*, vol. 32, 15 Suppl. 47 (1998) p. 15-21; and
 - Prasad: Acute and chronic angiotensin-1 receptor antagonism reverses endothelial dysfunction in atherosclerosis, *Circulation*, vol. 101 (2000), p. 2349 ff.
- 20 It is also known that ANG II antagonists selectively block the AT1 receptor, while the AT2 receptor which plays a part in anti-growth effects and tissue regeneration effects remains unaffected.
- EP-A-1 013 273 also describes the use of AT1-receptor antagonists or AT2-
- 25 receptor modulators for the treatment of diseases associated with an increase in the AT1-receptors in the sub-epithelial region or an increase in the AT2-receptors in the epithelium, particularly for the treatment of various lung diseases.
- In another aspect it was found that hypertension is often present at the same time
- 30 as hyperlipidaemia. Both symptoms are regarded as serious risk factors in the development of cardiovascular diseases, which often lead to unfavourable cardiovascular events.
- High blood cholesterol levels and high blood lipid levels are involved for example
- 35 in the start of atherosclerosis, a condition characterised by unevenly distributed

- 5 lipid deposits inside the arteries, including the coronary, carotid and peripheral arteries.

This irregular lipid distribution is thus characteristic of coronary heart damage, cardiovascular diseases, the gravity and prevalence of which are also affected by
10 the existence of diabetes, the sex of the person, cigarette smoking and left ventricular hypertrophy occurring as a side effect of hypertension (Wilson et al., Am. J. Cardiol., vol. 59(14) (1987), p. 91G-94G).

Type 2 diabetes mellitus is the manifestation of two pathophysiological
15 phenomena, namely a reduced secretion of insulin from the beta cells of the pancreas and insulin resistance in the target organs of the liver, skeletal musculature and fatty tissue. As a rule there is a complex disruption of both components. The disease is diagnosed as fasting hyperglycaemia, i.e. the blood sugar concentration after 10-12 hours' fasting is above the threshold of 125 mg of
20 glucose per dl of plasma. Controlled treatment of manifest type 2 diabetes can be achieved using compounds of the category of the thiazolidinediones (glitazones). These compounds improve the utilisation of circulating insulin and thus result in a lowering of the blood sugar levels (insulin sensitisers). At the same time the increased insulin levels are reduced by feedback mechanisms and in this way the
25 load on the pancreas is relieved. Insulin sensitisers such as troglitazone, rosiglitazone or pioglitazone develop this activity by binding to specific nuclear receptors known as PPAR-gamma (Peroxisomal Proliferator Activated Receptor).

WO 95/06410 discloses the use of angiotensin II receptor antagonists for treating
30 chronic inflammatory diseases including systemic autoimmune diseases.

Diabetes is mentioned as one of a number of examples of systemic autoimmune diseases. The autoimmune diseases include type 1 diabetes mellitus which occurs mainly in young people under 30 years of age with a genetic predisposition, in whom insulinitis occurs under the influence of various factors with subsequent

- 5 destruction of the B cells so that the pancreas can only produce a little insulin or none at all. Type 2 diabetes mellitus is not regarded as an autoimmune disease.

- Because every other type 2 diabetes patient shows signs of coronary heart disease at the time of diagnosis, for example, the causes of diabetes are
- 10 increasingly suspected to reside in a complex metabolic disorder which may be indicated by a number of risk factors such as abnormal glucose tolerance, increased fasting blood sugar, insulin resistance, high blood pressure, dyslipidaemia or centripetal obesity. The prevalence of insulin resistance is particularly marked in patients with hypertriglyceridaemia and low HDL (high-
- 15 density lipoprotein)-cholesterol. Reference is made to pre-type 2 diabetes, metabolic syndrome, syndrome X or insulin resistance syndrome. In a first phase a reduced insulin response by the target organs causes an increase in the pancreatic insulin secretion in order to keep the blood sugar level in the normal range. After a number of years of excessive or increasing insulin production there
- 20 comes a time when the insulin secretion by the beta cells of the pancreas cannot be increased any further. The phase of abnormal glucose tolerance then begins. The body can no longer absorb glucose peak values fast enough. Finally, if the fasting blood sugar remains persistently high, diabetes is manifest.
- 25 Angina pectoris, a condition characterised by severe constricting pains in the chest, often radiating out from the heart area to the left shoulder and down to the left arm, is frequently treated with a combination therapy of β -blockers, nitrate or calcium channel blockers together with a lipid lowering agent. Angina pectoris is often the result of cardiac ischaemia and is normally caused by coronary disease.
- 30 When treated surgically, angina patients often suffer complications such as restenosis which is experienced either as a short term proliferative reaction to the trauma caused by the angioplasty or as a long-term progression of the arteriosclerotic process both in transplanted vessels and in angioplasty segments. Some possible treatments for lowering lipids and cholesterol are based on
- 35 inhibiting the activity of the enzyme 3-hydroxy-3-methylglutaryl-coenzyme A-

5 reductase (HMG-CoA-reductase), which catalyses the conversion of 3-hydroxy-3-methylglutaryl-coenzyme A into mevalonate, an early stage in the biosynthetic cholesterol metabolic pathway. Known inhibitors of HMG-CoA-reductase are for example compounds derived from a fungal metabolite the names of which end in "statin", such as pravastatin, lovastatin, fluvastatin, simvastatin or atorvastatin.

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Atorvastatin is a potent inhibitor of the enzyme 3-hydroxy-3-methylglutaryl-coenzyme A-reductase (HMG-CoA-reductase) and is known as a high-grade liver-selective inhibitor of cholesterol biosynthesis, the effect of which involves lowering Low Density Lipoprotein Cholesterol (LDL-C). These activities are the reason for
15 the attractiveness of this molecule in the treatment of combined hyperlipidaemia, a normal atherogenic disorder in clinical practice, and thus also in preventing the progression of atheroma.

Investigations have also shown that lowering the LDL-C level provides protection
20 against coronary heart diseases (cf. for example "Scandinavian Simvastatin Survival Study" or 4S study, published in The Lancet, vol. 344 (1994), p. 1383-1389, or the study "Prevention of coronary heart disease with pravastatin in men with hypercholesterolemia", published by Shepherd et al., in The New England Journal of Medicine, vol. 333 (1995), p. 1301-1307).

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Other studies are being carried out to determine the protective effect of statins against the occurrence of heart attacks, strokes and coronary heart diseases in non-insulin-dependent diabetics; "Collaborative Atorvastatin Diabetes Study" or the CARDS study "Atorvastatin Versus Revascularisation Treatment", the AVERT
30 study and the "Anglo-Scandinavian Cardiac Outcomes trial" or ASCOT study.

Since high blood pressure often occurs together with hyperlipidaemia or signs of type 2 diabetes, as already mentioned, and since these signs are main risk factors for the development of cardiovascular diseases which often lead to unfavourable
35 cardiovascular events, it would be beneficial for the patient to have access to a

- 5 single therapy which prevents or treats these conditions. It would also be advantageous if the combination therapy also brought about an improvement in the prevention or treatment of cardiopulmonary, pulmonary or renal diseases for which ANG II antagonists have been found to be effective.
- 10 The aim of the present invention is to provide pharmaceutical compositions which are suitable both for the treatment of high blood pressure and also for the treatment of hyperlipidaemia and simultaneously for the treatment of manifest type 2 diabetes and also for the treatment of first indications of the complex metabolic disorder of prediabetes and hence may be used to prevent type 2 diabetes
- 15 mellitus.
- Combined treatments and corresponding compositions which contain HMG-CoA-reductase inhibitors and ANG II antagonists have already been proposed.
- WO-95/26188 describes a method of treating atherosclerosis and reducing
20 cholesterol, using an HMG-CoA-reductase-inhibitor and an ANG II antagonist. Pravastatin, simvastatin and lovastatin are mentioned as possible HMG-CoA-reductase inhibitors which may be used. Losartan is mentioned as an ANG II-antagonist which may possibly be used.
 - WO-97/37688 describes the combined use of HMG-CoA-reductase inhibitors
25 and ANG II antagonists for the treatment of numerous conditions, including hypertension and atherosclerosis. Pravastatin, simvastatin, lovastatin and fluvastatin are mentioned as possible HMG-CoA-reductase inhibitors which may be used.
 - WO-99/11260 describes the combined use of a special HMG-CoA-reductase-
30 inhibitor and ANG II antagonists for lowering blood pressure and the lipid levels and also for treating angina pectoris and atherosclerosis in mammals. The particular HMG-CoA-reductase-inhibitor is atorvastatin. Losartan, irbesartan and valsartan are mentioned as possible ANG II antagonists which are preferably used. Other ANG II antagonists mentioned are candesartan and
35 eprosartan.

- 5 • WO-00/45818 describes the combined use of an HMG-CoA-reductase-inhibitor and an ANG II antagonist for alleviating diabetic neuropathy and particularly for improving the conductive speed of the nerves and blood flow to the nerves in patients suffering from diabetes. The above examples of possible combinations are combinations comprising the statins pravastatin, simvastatin, cerivastatin, 10 fluvastatin, atorvastatin and statin (E) together with the ANG II antagonists losartan, irbesartan, valsartan and candesartan, of which candesartan is preferred.
- 15 • WO-01/15674 describes the combination of an inhibitor of the Renin-Angiotensin-System together with another antihypertensive, cholesterol-lowering agent, a diuretic or aspirin for preventing cardiovascular events such as stroke, congestive heart failure, cardiovascular death, myocardial infarct, worsening of angina, stoppage of the heart, revascularisation processes, diabetes and diabetic complications. Examples of possible combinations are the combinations of Angiotensin-Converting-Enzyme (ACE) inhibitors, i.e. 20 compounds whose names end in "-pril", such as captopril, imidapril, ramipril and the like, with the cholesterol level lowering agents lovastatin, pravastatin, simvastatin or fluvastatin.

Summary of the Invention

- 25 The present invention relates to a method for the prevention or treatment of cardiovascular, cardiopulmonary, pulmonary or renal diseases by improving endothelial function and achieving protection of organs, tissues and blood vessels in indications in which control of blood pressure and lipid levels are necessary, particularly in people in whom type 2 diabetes mellitus has been diagnosed or who 30 are suspected of prediabetes, for preventing diabetes and prediabetes, or for the treatment of Metabolic Syndrome and insulin resistance in patients with normal blood pressure. The process comprises generally administering effective amounts of telmisartan or a polymorph or salt thereof and atorvastatin. The invention further relates to suitable pharmaceutical compositions which contain telmisartan 35 or a polymorph or salt thereof and atorvastatin, as a combined preparation for

- 5 simultaneous, separate or sequential use in the prevention or treatment of these diseases.

Detailed Description

- 10 Within the scope of the present invention it has now surprisingly been found that the angiotensin II receptor antagonist telmisartan and the salts thereof not only act to reduce blood pressure, in known manner, but are also capable of increasing the expression of genes in a cellular system, the transcription of which is known to be regulated by the PPARgamma receptor. In order to ensure comparable conditions this effect is observed and quantified within the scope of the present invention by
- 15 means of a stably transformed cell line (cf. Example 2). The cells used are CHO cells which are the result of transformation with two gene constructs. The first of these constructs codes for the luciferase gene from *Photinus pyralis* (de Wet JR, Mol Cell Biol (1987) 7:725) under the control of a synthetic promoter with a five-fold repeat of a yeast Gal4 binding site (cf. GeneBank Sequence AF058756).
- 20 The second construct codes for a fusion protein consisting of the ligand binding domain of the human PPARgamma2 transcription factor (cf. GeneBank Sequence U79012) and the yeast GAL4 DNA binding domain (Amino acids 1-147; Sadowski I, Nucleic Acids Res (1989) 17:7539).
- 25 The induction of the transcription of PPARgamma-regulated genes is known from the thiazolidinediones used as antidiabetic drugs (e.g. rosiglitazone) and is brought about by their binding to the PPARgamma Receptor and its activation. Within the scope of the test system used here this effect may be quantified as an induced luciferase activity of the transformed cell line. In the case of telmisartan,
- 30 contrary to expectation, the same induction of a luciferase activity does not take place by the binding of the active substance to the PPARgamma Receptor. Binding of telmisartan to the PPARgamma receptor cannot be detected in various test systems. It is therefore presumed that the increase in the affinity of cofactor proteins for PPARgamma caused by the angiotensin II receptor antagonist
- 35 telmisartan also leads to the recruiting of the cofactor proteins if there are no high-

- 5 affinity synthetic PPARgamma ligands present. This then brings about activation of the transcription of genes regulated by the PPARgamma receptor, this activation being mediated by these cofactors. As the induction of these genes is responsible for the anti-diabetic activity of the thiazolidinediones it can be assumed that the induction of the same genes by telmisartan results in a
- 10 comparable anti-diabetic activity. Thus, these active substances are suitable not only for treating high blood pressure but also for treating and preventing type 2 diabetes mellitus. This includes the treatment and prevention of metabolic syndrome, syndrome X or insulin-resistance syndrome.
- 15 The discovery of this new therapeutic effect of telmisartan and the salts thereof means that they can be used to produce a pharmaceutical composition for the treatment of people or mammals in whom the prevention or treatment of cardiovascular, cardiopulmonary, pulmonary or renal diseases is indicated, particularly if type 2 diabetes mellitus has been diagnosed or if there is a suspicion
- 20 of prediabetes or who has been diagnosed with the metabolic disorder known as insulin resistance syndrome.

Of particular importance is the treatment of people in whom prevention or treatment of hypertension combined with hyperlipidaemia or atherosclerosis is

25 indicated, or the treatment of asthma, bronchitis or interstitial lung diseases. They are also suitable for the treatment and prevention of type 2 diabetes and pre-type 2 diabetes. This includes the treatment and prevention of Metabolic Syndrome, Syndrome X, or Insulin Resistance Syndrome. Of particular importance is the treatment of people in whom prevention or treatment of hypertension combined

30 with hyperlipidaemia or atherosclerosis is indicated, or the treatment of asthma, bronchitis or interstitial lung diseases.

Type 2 diabetes mellitus manifests itself in a fasting blood sugar level exceeding 125 mg of glucose per dl of plasma; the measurement of blood glucose values is a

35 standard procedure in routine medical analysis. If a glucose tolerance test is

- 5 carried out, the blood sugar level of a diabetic will be in excess of 200 mg of glucose per dl of plasma 2 hours after 75 g of glucose have been taken on an empty stomach. In a glucose tolerance test 75 g of glucose are administered orally to the patient being tested after 10-12 hours of fasting and the blood sugar level is recorded immediately before taking the glucose and 1 and 2 hours after
- 10 taking it. In a healthy subject the blood sugar level before taking the glucose will be between 60 and 110 mg per dl of plasma, less than 200 mg per dl 1 hour after taking the glucose and less than 140 mg per dl after 2 hours. If after 2 hours the value is between 140 and 200 mg this is regarded as abnormal glucose tolerance.
- 15 If insulin resistance can be detected this is a particularly strong indication of the presence of prediabetes. Thus, it may be that in order to maintain glucose homoeostasis one person needs 2-3 times as much insulin as another person, without this having any direct pathological significance. The most certain method of determining insulin resistance is the euglycaemic-hyperinsulinaemic clamp test.
- 20 The ratio of insulin to glucose is determined within the scope of a combined insulin-glucose infusion technique. There is found to be insulin resistance if the glucose absorption is below the 25th percentile of the background population investigated (WHO definition). Rather less laborious than the clamp test are so called minimal models in which, during an intravenous glucose tolerance test, the
- 25 insulin and glucose concentrations in the blood are measured at fixed time intervals and from these the insulin resistance is calculated. Another method of measurement is the mathematical HOMA model. The insulin resistance is calculated by means of the fasting plasma glucose and the fasting insulin concentration. In this method it is not possible to distinguish between hepatic and
- 30 peripheral insulin resistance. These processes are not really suitable for evaluating insulin resistance in daily practice. As a rule, other parameters are used in everyday clinical practice to assess insulin resistance. Preferably, the patient's triglyceride concentration is used, as increased triglyceride levels correlate significantly with the presence of insulin resistance.

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5 Thus, there is a suspicion of prediabetes if the fasting blood sugar level is above
the normal maximum level of 110 mg of glucose per dl of plasma but does not
exceed the threshold of 125 mg of glucose per dl of plasma which indicates
diabetes. Another indication of prediabetes is abnormal glucose tolerance, i.e. a
10 blood sugar level of 140-200mg of glucose per dl of plasma 2 hours after taking 75
g of glucose after a fast within the scope of a glucose tolerance test.

A triglyceride blood level of more than 150 mg/dl also indicates the presence of
pre-diabetes. This suspicion is confirmed by a low blood level for HDL cholesterol.
In women, levels below 40 mg per dl of plasma are regarded as too low while in
15 men levels below 50 mg per dl of plasma are regarded as too low. Triglycerides
and HDL cholesterol in the blood can also be determined by standard methods in
medical analysis and are described for example in Thomas L (Editor): "Labor und
Diagnose", TH-Books Verlagsgesellschaft mbH, Frankfurt/Main, 2000. A
suspicion of prediabetes is further confirmed if the fasting blood sugar levels also
20 exceed 110 mg of glucose per dl of plasma. If the blood levels measured are in
the region of these threshold values, the ratio of the waist measurement to the hip
measurement can be used as an additional aid to make the decision. If this ratio
exceeds a value of 0.8 in women or 1 in men, treatment is indicated.

25 Telmisartan is particularly indicated for treating diabetes or suspected prediabetes
if hypertension also has to be treated. This is the case if the systolic blood
pressure exceeds a value of 140 mm Hg and diastolic blood pressure exceeds a
value of 90 mm Hg. If a patient is suffering from manifest diabetes it is currently
recommended that the systolic blood pressure be reduced to a level below
30 130 mm Hg and the diastolic blood pressure be lowered to below 80 mm Hg. To
achieve these levels it may be indicated in certain cases to combine angiotensin II
receptor antagonists with a diuretic or a calcium antagonist. The term "diuretic"
includes thiazides or thiazide analogues such as hydrochlorothiazides (HCTZ),
clopamide, xipamide or chlorthalidone, aldosterone antagonists such as
35 spironolactone or eplerenone and also other diuretics suitable for treating high

- 5 blood pressure such as furosemide and piretanide, and combinations thereof with
amiloride and triamterene.

The present invention means that for subjects being treated for increased blood
pressure, the angiotensin II receptor antagonist telmisartan is indicated whenever
10 the development of prediabetes is to be prevented or manifest diabetes is to be
treated.

In only 10% of all cases of elevated blood pressure (secondary hypertension) is it
possible to determine an identifiable cause such as e.g. kidney disease. As a rule,
15 secondary hypertension can be remedied by treating and removing the cause.
However, in almost 90% of all cases it is primary hypertension, the exact cause of
which is not known and which therefore cannot be directly cured. The negative
effects of elevated blood pressure can be reduced by changing lifestyle and
correct treatment. The interaction of different risk factors or the combined
20 occurrence of individual risk factors appear to cause high blood pressure. In
particular, the combination of high blood pressure with disorders of the fat and
sugar metabolism is observed to an increasing extent. These disorders are often
unnoticed to begin with but can be recognised from increased blood levels of
triglycerides and glucose and lower blood levels of HDL cholesterol. At a fairly
25 advanced stage they can also be detected in slowly increasing corpulence. These
disorders can be explained by increasing insulin resistance. The less effective the
insulin, the more the fat and sugar metabolisms are disrupted. The combination of
all these disorders in the last analysis increases the probability of contracting the
sugar disease diabetes and dying prematurely of heart or vascular disease.

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As primary or essential hypertension is a multifactorial disease, it seems unlikely
that insulin resistance or hyperinsulinaemia is the sole cause of high blood
pressure. A number of observations indicate, however, that defects in the insulin
metabolism have a hypertensive effect and thus predispose to high blood
35 pressure. In connection with this, reference may be made to hypertensive insulin

5 resistance. Thus the presence of insulin resistance can be detected in about 50% of normal-weight hypertensives and normotensive close relatives. In obese patients not only is there a higher level of insulin resistance, but also a stronger correlation between hypertension and hyperinsulinaemia than in slim hypertensives.

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Estimates are based on the supposition that about a third of adults in those parts of the world with an excessive supply of food are affected by the combination of high blood pressure and disorders of the fat and sugar metabolism and that this number will continue to increase. Consequently there is a need for drugs which
15 are capable of helping to slow down or stop the progress of the above-mentioned metabolic disorders at the earliest possible stage and at the same time to obviate the detrimental effects of increased blood pressure on the health.

The present invention now discloses a pharmaceutical composition which can be
20 used both to treat hypertension and hyperlipidaemia simultaneously and to treat manifest type 2 diabetes or the first signs of the complex metabolic disorder of prediabetes. The new active substance combination is particularly suitable for the treatment and prevention of the above-mentioned hypertensive insulin resistance, which denotes insufficient utilisation of the insulin circulating in the bloodstream,
25 combined with a resulting increase in blood pressure. Thus, the invention also includes diabetes prevention in patients who are being treated for high blood pressure and hyperlipidaemia. If therefore the combination of telmisartan and atorvastatin is used immediately to control blood pressure, hyperlipidaemia or
hypertensive insulin resistance as soon as one of the above-mentioned signs of
30 prediabetes is present, the onset of manifest type 2 diabetes can be delayed or prevented.

5 Telmisartan and the suitable salts thereof thus:

- do not exhibit any *in vitro* binding to the ligand binding domain of a human PPARgamma receptor, but lead to the
- induction of a luciferase activity when they are added to the culture medium of a stably transformed PPARgamma reporter cell line which

- 10 a) expresses a fusion protein consisting of the ligand binding domain of the human PPARgamma transcription factor and the yeast GAL4 DNA binding domain and
- b) a luciferase gene under the control of a five-times repeated yeast Gal4 binding site.

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The preparation of a PPARgamma reporter cell line of this kind is described in Example 2.

There is no *in vitro* binding to the ligand binding domain of the human

- 20 PPARgamma2 receptor if it cannot be detected in an AlphaScreen (Ullmann EF et al, Proc Natl Acad Sci USA (1994) 91:5426-5430). Instead of an Alpha Screen, an SPA assay (Mukherjee R et al., J Steroid Biochem Mol Biol (2002) 81:217-225) or an NMR investigation (Johnson BA et al., J Mol Biol (2000) 298:187-194) may also be carried out. As a rule, binding to the receptor cannot be detected by any of
- 25 these methods.

- If it appears useful or necessary to use an angiotensin II receptor blocker in conjunction with one or more other therapeutic active substances, telmisartan is a preferred angiotensin II receptor blocker, as it combines a blood pressure lowering and antidiabetic activity in a single active substance and helps to prevent diabetes.
- 30 For this reason, preformulated active substance combinations of telmisartan with the HMG-Co A reductase inhibitor atorvastatin constitute a major further development in the treatment of cardiovascular, cardiopulmonary, pulmonary or renal diseases, but particularly when there is a need to treat hyperlipidaemia,

- 5 prediabetes or manifest type 2 diabetes, osteoporosis or Alzheimer's simultaneously, as well as prevent diabetes.

It is observed that by joint administration of an effective amount of telmisartan with an effective amount of atorvastatin, or polymorphs or salts thereof, surprising
10 advantages can be achieved in the prevention or treatment of cardiovascular, cardiopulmonary, pulmonary or renal diseases in patients requiring treatment with a high degree of efficacy, irrespective of the known hypotensive effect of the active substance telmisartan and independently of the antihyperlipidaemic activity of the active substance atorvastatin, compared with administering the ANG II antagonist
15 or the HMG-CoA-reductase inhibitor on its own. Thus, it is possible for example to control the expression of the Matrix Metalloproteinase MMP-9, which is expressed to a greater extent in chronic inflammation of the respiratory tract or in type 2 diabetes. Elevated plasma levels of the inflammation-promoting cytokine CD40L can also be counteracted. Increased plasma levels of CD40L are a known risk
20 factor for cardiovascular diseases.

It is also observed that the prevention or treatment improves endothelial function and affords protection of organs, tissues and blood vessels in diseases in which there is a need to control both blood pressure and also the lipid levels. Thus, the
25 elasticity of the arteries can be improved and in the skin an enhanced production of NO, a marker of endothelial function, can be achieved.

It is also observed that the prevention or treatment is particularly effective in the following situations:

- 30 • indications (A) which can be positively influenced by inhibition of the activities mediated by the AT1-receptor and maintenance of the activities of angiotensin II (ANG II) mediated by the AT2-receptor and by inhibition of the HMG-CoA-reductase activities, by means of which the activities mediated by bradykinin can thus be potentiated and antihyperlipidaemic activities can be achieved; or

- 5 • indications (B) which go hand in hand with an increase in the AT1 receptors in the subepithelial region or an increase in the AT2 receptors in the epithelium.

Suitable indications (A) are selected from the following indications:

- treatment of combined hypertension and hyperlipidaemia;
- 10 • reduced occurrence of stroke, acute myocardial infarct or cardiovascular deaths, particularly in people with an increased risk of adverse cardiovascular events or strokes;
- provision of a renoprotective effect, e.g. in renal failure or diabetic nephropathy;
- prevention of left ventricular hypertrophy, vascular thickening, e.g. prevention
- 15 of the thickening of blood vessel walls after vascular surgery, improvement of the chances of survival after heart transplants, prevention of arterial restenosis after angioplasty, prevention or treatment of atherogenic disorders such as atherosclerosis, protection against coronary artery diseases, prevention of atheroma progression and prevention of diabetic angiopathy;
- 20 • lowering of cholesterol, lowering of plasma-fibrinogen and plasma viscosity, inhibition of the proliferation of smooth muscle cells, reduction of the ability of macrophages to oxidise LDL, protection of heart muscle cells from hypoxic damage and lowering of the plasminogen activator inhibitor 1 (PAI-1);
- prevention or treatment of ischaemic peripheral circulatory disorders and
- 25 myocardial ischaemia (angina); and
- prevention of the progression of heart failure after myocardial infarct.

- Suitable indications (B) are selected from the following indications: obstructive respiratory diseases, chronic obstructive lung diseases such as bronchitis or
- 30 chronic bronchitis, emphysema, for example caused by asthma, cystic fibrosis, interstitial lung disease, lung cancer, pulmonary vascular diseases and increased resistance to the airflow in forced ventilation; adult respiratory distress syndrome (ARDS), reduction in the proliferative capacity of the epithelium in cancer of the lung and breast, treatment of sepsis syndromes, lung damage such as
- 35 inflammation of the lung, aspiration of the stomach contents, trauma to the

5 ribcage, shock, burns, fatty embolisms, heart-lung bypass, O₂ toxicity,
haemorrhagic pancreatitis, interstitial and bronchoalveolar inflammation,
particularly when accompanied by increased expression of Matrix
Metalloproteinase such as MMP-9, proliferation of epithelial and interstitial cells,
collagen accumulation and fibrosis.

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Thus, the present invention provides a process for the prevention or treatment of
hypertension and hyperlipidaemia, particularly in a mammal in whom diabetes has
been diagnosed or there is a suspicion of prediabetes, the process comprising the
combined administration of an effective amount of the HMG-CoA-reductase-
15 inhibitor atorvastatin or a polymorph or salt thereof, together with an effective
amount of the ANG II antagonist telmisartan or a polymorph or salt thereof.

The invention further relates to the combined use of telmisartan and atorvastatin or
the combined use of polymorphs or salts of these active substances in the
20 manufacture of a pharmaceutical composition for the prevention or treatment of
hypertension in combination with hyperlipidaemia, particularly if diabetes has been
diagnosed or there is a suspicion of prediabetes.

Thus, the advantageous activity of the processes according to the invention is
25 based primarily on the protective effective of the combined treatment for organs,
tissues and blood vessels, as well as the preventive effect in relation to diabetes.

The above-mentioned unexpected advantages may be attributable to a more
effective blockade of the activities of ANG II mediated by the AT1 receptor, to the
30 activity of ANG II mediated by the AT2 receptor, which remains unaffected by this
specific ANG II antagonist, together with an increase in the activities mediated by
bradykinin, to the PPARgamma-like transcription activation and to the
achievement of an antihyperlipidaemic activity by atorvastatin.

5 It is observed, for example, that the combined administration of the specific ANG II antagonist telmisartan with the specific HMG-CoA-reductase-inhibitor atorvastatin or the combined administration of polymorphs or salts of these active substances brings about a significant prevention of cardiovascular deaths and overall mortality, particularly in respect of the occurrence of stroke and acute myocardial
10 infarct, compared with the administration of one of these active substances on its own.

Therefore, a preferred process according to the invention comprises reducing the occurrence of stroke and acute myocardial infarct in people or non-human
15 mammals requiring treatment, particularly in individuals with manifest type 2 diabetes or suspected prediabetes or with a increased risk of adverse cardiovascular events or stroke, by administering telmisartan together with atorvastatin or by administering polymorphs or salts of these active substances together.

20 It is observed, moreover, that the combined treatment and the corresponding compositions which specifically contain an amount of the HMG-CoA-reductase inhibitor atorvastatin together with an amount of the ANG II antagonist telmisartan or polymorphs or salts of these active substances, result in a high activity in the
25 regulation of blood pressure and in lipid regulation in mammals. It is expected that the synergistic activity achieved using this special combination is surprisingly superior to the activity of corresponding conventional combinations.

By a synergistic combination for regulating blood pressure and lipids is meant that
30 it contains an amount of atorvastatin and an amount of telmisartan or polymorphs or salts of these active substances, wherein the quantity of the individual active substance is not sufficient on its own to achieve the therapeutic effect which is obtained by administering the combination of agents, and the combined effects of the quantities of therapeutic agents are greater than the sum of the therapeutic

- 5 activities which can be achieved with the quantities of the individual therapeutic agents.

The present invention further relates to pharmaceutical compositions containing telmisartan or one of the salts thereof combined with atorvastatin and the
10 preparation thereof. The pharmaceutical compositions are used for treating human or non-human mammals for the prevention or treatment of the above-mentioned diseases or indications and contain telmisartan and atorvastatin, optionally together with pharmaceutically acceptable diluents and/or carriers, in the form of a combined preparation for simultaneous, separate or successive use in
15 the prevention or treatment of these diseases or indications.

These combinations of active substances are generally incorporated with one or more formulation adjuvants such as mannitol, sorbitol, xylitol, saccharose, calcium carbonate, calcium phosphate, lactose, croscarmellose sodium salt (cellulose
20 carboxymethylether sodium salt, cross-linked), crospovidone, sodium starch glycolate, hydroxypropylcellulose (low-substituted), maize starch, polyvinylpyrrolidone, copolymers of vinylpyrrolidone with other vinyl derivatives (copovidone), hydroxypropylcellulose, hydroxypropylmethylcellulose, microcrystalline cellulose or starch, magnesium stearate, sodium stearyl fumarate,
25 talc, hydroxypropylmethylcellulose, carboxymethylcellulose, cellulose acetate phthalate, polyvinyl acetate, water, water/ethanol, water/glycerol, water/sorbitol, water/polyethyleneglycol, propyleneglycol, cetylstearyl alcohol, carboxymethylcellulose or fatty substances such as hard fat or suitable mixtures thereof, into conventional galenic preparations such as plain or coated tablets,
30 capsules, powders, suspensions or suppositories.

Tablets may be obtained for example by mixing the active substance or substances with one or more excipients and subsequently compressing them. The tablets may also consist of several layers.
35

5 Examples of excipients are:

- inert diluents such as mannitol, sorbitol, xylitol, saccharose, calcium carbonate, calcium phosphate and lactose;
- disintegrants such as croscarmellose sodium salt (cellulose carboxymethylether sodium salt, cross-linked), crospovidone, sodium starch glycolate, hydroxypropylcellulose (low-substituted) and maize starch;
- 10 • binders such as polyvinylpyrrolidone, copolymers of vinylpyrrolidone with other vinyl derivatives (copovidone), hydroxypropylcellulose, hydroxypropylmethylcellulose, microcrystalline cellulose or starch;
- lubricants such as magnesium stearate, sodium stearyl fumarate and talc;
- 15 • agents for achieving delayed release such as hydroxypropylmethylcellulose, carboxymethylcellulose, cellulose acetate phthalate and polyvinyl acetate; and
- pharmaceutically permitted colourings such as coloured iron oxides.

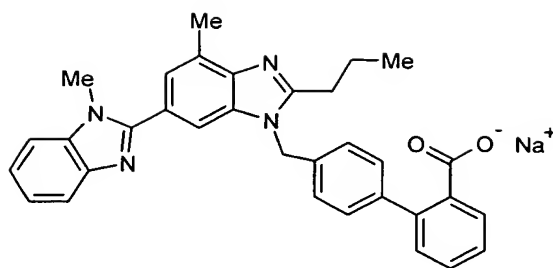
In all aspects of the present invention the particular ANG II antagonist telmisartan is {4'-[2-n-propyl-4-methyl-6-(1-methylbenzimidazol-2-yl)-benzimidazol-1-ylmethyl]-biphenyl-2-carboxylic acid} or polymorphs or salts thereof, preferably the sodium salt. Telmisartan is already on the market, e.g. under the name Micardis®.

Telmisartan is described for example in EP-0 502 314 and US-5 591 762.

25 Polymorphs of telmisartan are described for example in WO-00/43370, US-6 358 986 and US-6 410 742. Sodium salts of telmisartan are described for example in WO 03/037876.

For example it states in WO 03/037876 that the sodium salt of telmisartan of

30 formula:



5

can be selectively obtained in a crystalline polymorphic form by a suitable choice of the manufacturing conditions.

- 10 This crystalline form of the sodium salt of telmisartan is characterised by the melting point $T = 245 \pm 5$ °C (determined by differential scanning calorimetry using the Mettler-Toledo DSC82 apparatus; heating rate: 10° K/min).

- 15 The sodium salt of telmisartan may be prepared using one of the following two manufacturing processes.

- According to all aspects of the invention the HMG-CoA-reductase inhibitor is atorvastatin or polymorphs or salts thereof, preferably the hemicalcium salt {[R-(R*,R*)]-2-(4-fluorophenyl)- β,δ -dihydroxy-5-(1-methylethyl)-3-phenyl-4-
20 [(phenylamino)-carbonyl]-1H-pyrrol-1-heptanoic acid hemicalcium salt), which is marketed for example under the brand names Lipitor®, Zarator® and Sortis®.

- Atorvastatin is described for example in EP 0 247 633 and US-4 681 893. Polymorphs of atorvastatin are described for example in WO-97/03958, WO-
25 97/03959, EP-0 848 704 and EP-1 148 049. Salts of atorvastatin (monopotassium salt, monosodium salt, calcium salt, magnesium salt, zinc salt and meglumine) are described for example in EP-0 409 281 and US-5 273 995.

- 30 By combined administration of the two active substances is meant a successive or simultaneous administration, of which simultaneous administration is preferred.

- 5 For successive administration telmisartan may be given before or after the administration of atorvastatin.

The active substances may be administered by oral, buccal or parenteral route, by inhalation, or rectally or topically; oral administration is preferred. Parenteral
10 administration may comprise subcutaneous, intravenous, intramuscular and intrasternal injections as well as infusion techniques.

The active substances may be given orally in a variety of different dosage forms, i.e. they may be prepared with various pharmaceutically acceptable inert carriers
15 to form tablets, capsules, pastilles, sweets, powders, sprays, aqueous suspensions, elixirs, syrups and the like. Such carriers include solid diluents or fillers, sterile aqueous media and various non-toxic organic solvents. In addition, oral pharmaceutical preparations of this kind may be provided with suitable
20 sweeteners and/or flavourings, using various agents conventionally used for such purposes. In general the compounds according to the invention are present in oral formulations of this kind in concentrations ranging from about 0.5 to about 90 wt%, based on the total composition, in amounts such that they produce the desired dosage units. Other suitable dosage forms for the compounds according to the
25 invention include preparations and devices with controlled release, with which the skilled man will be familiar.

For oral administration it is possible to use tablets containing various carriers such as sodium citrate, calcium carbonate and calcium phosphate together with various disintegrants, such as starch and preferably potato or tapioca starch, alginic acid
30 and certain complex silicates, together with binders such as polyvinylpyrrolidone, saccharose, gelatine and gum arabic. Lubricants such as magnesium stearate, sodium laurylsulphate and talc or compositions of a similar type may also be used as fillers in filled soft and hard gelatine capsules. These may also include lactose or milk sugar as well as high molecular polyethyleneglycols. If aqueous
35 suspensions and/or elixirs are desired for oral administration, the active

- 5 substances may be combined with various sweeteners or flavourings, colouring agents or dyes and optionally emulsifiers and/or water, ethanol, propyleneglycol, glycerol and various combinations thereof.

For parenteral administration solutions of the compounds in sesame or groundnut
10 oil or in aqueous propyleneglycol as well as sterile aqueous solutions of the corresponding pharmaceutically acceptable salts may be used. Aqueous solutions of this kind may optionally be suitably buffered and the liquid diluent may optionally be made isotonic with sufficient quantities of common salt or glucose. These special aqueous solutions are particularly suitable for intravenous, intramuscular
15 and subcutaneous injection. Sterile aqueous media may easily be obtained by conventional methods known to the skilled man. For example, distilled water is normally used as a liquid diluent. The finished preparation is passed through a suitable bacterial filter, e.g. a filter made of sintered glass, diatomaceous earth or unglazed porcelain. Preferred filters of this type include Berkefeld, Chamberland
20 and asbestos disc metal Seitz filters, the liquid being aspirated into a sterile container using a suction pump. Throughout the entire process of preparing these injectable solutions the necessary steps should be carried out in such a way as to obtain the end products in a sterile state.

- 25 For transdermal administration the formulations of the special compounds or combinations include for example solutions, lotions, ointments, creams, gels, suppositories, delayed-release preparations and devices therefor. These formulations comprise the compound(s) in particular and may contain ethanol, water, penetration promoters and inert carriers, e.g. gel-forming materials, mineral
30 oil, emulsifiers, benzyl alcohol and the like.

The formulations prepared contain, for example, an equivalent of 2.5-40 mg, preferably 5, 10, 15, 20, 25, 30, 35 or 40 mg of atorvastatin. Atorvastatin or polymorphs or salts thereof may be administered in daily doses of about 1.25 mg
35 (or 0.018 mg/kg of body weight, based on a person weighing 70 kg) to about

- 5 450 mg (6.43 mg/kg of body weight, based on a person weighing 70 kg) by oral route, about 20 mg (0.286 mg/kg of body weight, based on a person weighing 70 kg) by parenteral route and preferably in a dosage of about 2.5 mg (0.036 mg/kg of body weight, based on a person weighing 70 kg) to about 80 mg (1.428 mg/kg of body weight, based on a person weighing 70 kg) by oral route.
- 10 Particularly preferred is an oral daily dose of about 5 mg (0.071 mg/kg of body weight, based on a person weighing 70 kg), about 10 mg (0.143 mg/kg of body weight, based on a person weighing 70 kg), about 20 mg (0.286 mg/kg of body weight, based on a person weighing 70 kg) or about 40 mg (0.571 mg/kg of body weight, based on a person weighing 70 kg) or, especially to start with, an oral daily
- 15 dose of about 10 mg by oral route.

The formulations prepared contain, for example, an equivalent of 20-200 mg, preferably 20, 40, 80, 120, 160 or 200 mg of the free acid of telmisartan. If the active substance is combined with hydrochlorothiazide (HCTZ) or clorthalidone,

20 the formulation contains 10-50 mg, preferably 50, 25 or 12.5 mg of the diuretic. Telmisartan or polymorphs or salts thereof may be administered in a daily dose of 10 mg (or 0.143 mg/kg of body weight, based on a person weighing 70 kg) to 500 mg (7.143 mg/kg of body weight, based on a person weighing 70 kg) by oral route and about 20 mg (0.286 mg/kg of body weight, based on a person weighing 70 kg)

25 by parenteral route, preferably 20 mg (0.286 mg/kg of body weight, based on a person weighing 70 kg) to 100 mg (1.429 mg/kg of body weight, based on a person weighing 70 kg) by oral route. Particularly preferred is an oral daily dose of 40 mg (0.571 mg/kg of body weight, based on a person weighing 70 kg) to 80 mg (1.143 mg/kg of body weight, based on a person weighing 70 kg) or in particular a

30 dose of about 80 mg (1.143 mg/kg of body weight, based on a person weighing 70 kg).

Preferably the ratio of atorvastatin to telmisartan or the polymorphs or salts thereof in the pharmaceutical combination is 1:100 to 100:1 (based on weight).

- 5 In particularly preferred embodiments atorvastatin or a polymorph or salt thereof together with telmisartan or a polymorph or salt thereof is administered by oral route in the following daily doses:
- 10 mg of atorvastatin and 40 mg of telmisartan (or polymorphs or salts thereof);
 - 10 mg of atorvastatin and 80 mg of telmisartan (or polymorphs or salts thereof);
 - 10 20 mg of atorvastatin and 40 mg of telmisartan (or polymorphs or salts thereof);
 - 20 mg of atorvastatin and 80 mg of telmisartan (or polymorphs or salts thereof).

According to a preferred embodiment the pharmaceutical compositions according to the invention contain the HMG-CoA-reductase-inhibitor in an amount of 1.25 mg
15 to 450 mg and the ANG II antagonist in an amount of 10 mg to 500 mg in individual dosage units, optionally together with one or more pharmaceutically acceptable diluents and/or carriers.

According to another preferred embodiment the pharmaceutical compositions
20 according to the invention contain atorvastatin in an amount of 2.5 mg to 80 mg and telmisartan in an amount of 20 to 100 mg in individual dosage units, optionally together with one or more pharmaceutically acceptable diluents and/or carriers.

Another preferred sub-group of pharmaceutical compositions according to the
25 invention contain atorvastatin in an amount of 5 mg to 20 mg and telmisartan in an amount of 40 mg to 80 mg in individual dosage units, optionally together with one or more pharmaceutically acceptable diluents and/or carriers.

Another preferred sub-group of pharmaceutical compositions according to the
30 invention contain atorvastatin in an amount of 10 or 20 mg and telmisartan in an amount of 40 or 80 mg in individual dosage units, optionally together with one or more pharmaceutically acceptable diluents and/or carriers.

As already mentioned, the present invention also relates to the use of telmisartan
35 for preparing a pharmaceutical composition for treating the human or non-human

- 5 mammalian body for the prevention or treatment of the above-mentioned indications when used in combination with atorvastatin. By this use is meant the preparation of all the above-mentioned pharmaceutical compositions according to the invention.

10 Examples:

Example 1: Telmisartan, losartan and irbesartan do not bind *in vitro* to the PPARgamma ligand binding domain

- 15 Protein containing the human PPARgamma-ligand binding domain (LBD) was prepared as a GST fusion protein in *E.coli* and purified by affinity chromatography. To do this, a DNA section which codes for the amino acids 205-505 of the human PPARgamma2 transcription factor (cf. Genbank entry U79012) was subcloned via the additionally inserted restriction cutting sites BamH I and Xho I into the
- 20 expression vector pGEX-4T-1 (Amersham) and the sequence of the section was monitored. The fusion protein was expressed in the *E.coli* strain BL21(DE3) recommended for pGEX vectors after induction with 0.2mM IPTG for 4 hours at 25°C. The bacteria were pelleted after the induction and frozen in batches in PBS, pH 7.4. After opening up in a French Press, the dissolved GST-PPARgamma-
- 25 LBD-fusion protein was purified using a GSTrap column (Pharmacia). Elution was carried out by the addition of 20mM reduced glutathione. The GST-PPARgamma-LBD-protein fractions were desalinated using a HiTrap desalting column (Pharmacia) and the protein concentration was determined using a standard
- assay.

- 30 Protein containing the human RXRalpha ligand binding domain (LBD) was prepared as a His tag fusion protein in *E.coli* and purified by affinity chromatography. To do this a DNA section which codes for the amino acids 220-461 of the human RXRalpha transcription factor (cf. Genbank entry NM_002957,
- 35 nt 729-1457) was subcloned via the additionally introduced restriction cutting sites

5 BamH I and Not I into the expression vector pET28c (Novagen) and the sequence of the section was monitored. The fusion protein was expressed in the *E.coli* strain BL21(DE3) recommended for pET vectors after induction with 0.2mM IPTG for 4 hours at 25°C. The bacteria were pelleted after the expression and frozen in batches in PBS, pH 7.4. After opening up in a French Press, the dissolved His-
10 RXRalpha -LBD-fusion protein is purified using a HiTrap chelating column (Pharmacia). Elution was carried out using a 500 mM imidazole step. The His-RXRalpha -LBD protein fractions were desalinated using a HiTrap desalting column (Pharmacia) and the protein concentration was determined using a standard assay.

15

a) AlphaScreen

Alpha Screen assays were first described in Ullmann EF et al., Proc Natl Acad Sci USA (1994) 91:5426-5430. The measurements carried out within the scope of this Example were carried out as described by Glickman JF et al., J Biomol Screen
20 (2002) 7:3-10. The assay buffer consisted of 25mM Hepes pH7.4, 100mM NaCl, 1mM DTT, 0.1% Tween-20, 0.1% BSA. 3nM GST-PPARgamma-LBD fusion protein, 15nM biotinylated LXXLL peptide of the cofactor CBP (corresponding to the peptide disclosed on page 218 of Mukherjee R et al., J Steroid Biochem Mol Biol (2002) 81:217-225 with an additional N-terminal cysteine), and in each case
25 10µg/ml of anti-GST-acceptor beads or streptavidine donor beads (Applied Biosystems) were incubated in a total volume of 12.5µl in the presence of different concentrations of a test substance (in DMSO) for 4 hours at ambient temperature. The final DMSO concentration in the assay was 1% (v/v). A 1% DMSO solution was used as the background control (NSB). The measurement was done using a
30 Packard fusion measuring device.

conc. / M	telmisartan		rosiglitazon	
	MW	SD	MW	SD
NSB	619	21	573	17
1.00E-08			820	18
3.00E-08	642	41	1720	48
1.00E-07	606	10	8704	59
3.00E-07	644	56	27176	1232
1.00E-06	677	14	43233	1083
3.00E-06	720	35	52691	3771
1.00E-05	847	82	56366	4303
5.00E-05	1111	135		

5

Unlike rosiglitazone, a PPARgamma-agonist known from the literature with binding in the LBD, the use of increasing concentrations of telmisartan, losartan and irbesartan (concentrations of up to 50µM) did not result in any direct activation of the PPARgamma-LBD and hence in any significant recruiting of the LXXLL peptide.

10

b) SPA Assay

A description of the SPA assay format can be found in [Mukherjee R et al., J Steroid Biochem Mol Biol \(2002\) 81:217-225](#). The assay buffer consisted of 20mM Tris pH 7.5, 25mM KCl, 10mM DTT, 0.2% Triton X-100). Thirty (30)nM GST-PPARgamma-LBD fusion protein, 30nM His-RXRalpha-LBD, anti-GST-antibody (1:600, Amersham Pharmacia), 0.25mg protein A SPA PVT antibody-binding beads (Amersham Pharmacia), 30nM ³H-labelled rosiglitazone were incubated with dilutions of the test substance for 5 hours at room temperature in a total volume of 100µl.

15

20

- 5 10µM of unlabelled rosiglitazone was added as background control (NSB) instead of the radioactive rosiglitazone, and the solvent used, e.g. DMSO, was added as the maximum value (Bmax) instead of a test substance.

After the incubation the test preparations were centrifuged for 5 minutes at 2000

- 10 rpm in a Hettich Universal 30Rf centrifuge and measured using a Packard TopCount NXT.

conc / M	telmisartan		irbesartan		losartan	
	MW	SD	MW	SD	MW	SD
NSB	217	9	217	9	217	9
Bmax	911	15	911	15	911	15
1.00E-07	837	49	913	54	915	43
3.00E-07	802	28	810	49	835	11
1.00E-06	818	27	815	51	901	10
3.00E-06	818	20	779	26	814	53
1.00E-05	703	30	723	37	787	46
3.00E-05	691	222	648	40	784	96
1.00E-04	545	18	510	81	611	17

- 15 In contrast to direct PPARgamma-agonists which bind to the PPARgamma-LBD, no concentration-dependent displacement of the radioactive rosiglitazone from the binding pocket took place even in the presence of very large excesses of telmisartan, losartan or irbesartan.

5 c) NMR investigations

In contrast to a direct PPARgamma ligand, e.g. rosiglitazone, no interaction of the test substance with amino acids in the binding pocket takes place during the measurement of the ¹⁵N TROSY spectrum of the PPARgamma-LBD in the presence of the test substance telmisartan. The amino acids of the binding pocket
10 have the same position in the presence of the test substances as in the absence of a ligand.

15 Example 2: **Preparation of a stably transformed PPARgamma reporter cell line**

A DNA section which codes for amino acids 205-505 of the human PPARgamma2 transcription factor (corresponding to nucleotides 703-1605 of Genbank sequence U79012) was incorporated into the Multiple Cloning Site of the vector pFA-CMV
20 (Stratagene) via additionally inserted BamH I and Hind III restriction cutting sites and the sequence was verified. The resulting plasmid pFA-CMV/hPPARgamma2-LBD codes N-terminally of the PPARgamma-LBD in the same reading frame for a Gal4 DNA binding domain. In addition the plasmid codes for neomycin resistance.

25 The cell line CHO-K1 (ATCC CCL-61) was cotransfected with the plasmids pFA-CMV/hPPARgamma2-LBD and pFR-Luc (Stratagene). pFR-Luc codes for the luciferase gene under the control of a five-times repeated yeast Gal4 binding site. The transfection was carried out with lipofectamine2000 in accordance with the manufacturer's instructions.

30 After transfection the cells were cultivated in medium (Ham's F12 with 10% foetal calf serum) in the presence of 0.5 mg/ml G-418. After six days' cultivation the cells were passaged and kept in culture for another 10 days. The resulting neomycin-resistant colonies were picked out under the microscope and transferred
35 into 96 well dishes and cultured. Various transformed cell lines were obtained with

- 5 the plasmids contained therein (e.g. clone no. 10, 11, 13 etc), which were kept in the culture medium.

The cell lines were examined for the inducibility of the luciferase gene using a PPARgamma agonist, e.g. rosiglitazone, and react with an increased luciferase
10 signal to stimulation by the PPARgamma agonist.

Example 3: Telmisartan, losartan and irbesartan activate PPARgamma at cellular level

15

The CHO-K1 cell line derived from the transformed clone 11 of Example 2 was seeded in 96-well flat-bottomed dishes in a density of 3×10^4 cells/200µl/well and cultivated overnight in Ham's F-12 medium with 10% foetal calf serum and 0.5mg/ml G-418. After 24 hours the medium is changed for one without any
20 added G-418.

The test substances were brought to 100 times the desired concentration with a suitable solvent, e.g. DMSO, and diluted 1:100 with the medium placed in the cell culture plate. The solvent used, e.g. DMSO, was used as the background control
25 in the same concentration.

24 hours after the addition of the substance the supernatants were discarded and the cells were washed twice with 150µl washing buffer (25mM Tricine, 16.3mM MgSO₄, pH7.8). After the washing steps 50µl of washing buffer with 150µl of
30 luciferase assay buffer (25mM Tricine, 0.5mM EDTA, 0.54mM NaTPP, 16.3mM MgSO₄, 1.2mM ATP, 0.05mM luciferine, 56.8mM 2-mercaptoethanol, 0.1% Triton X-100, pH7.8) was added to each test preparation. Luminescence was measured after a five minute wait using a Packard TopCount NXT. The luciferase activity was obtained by integrating the relative luciferase units (RLU) of the first ten
35 seconds after the start of measurement.

conc / M	telmisartan		irbesartan		losartan		rosiglitazone	
	MW	SD	MW	SD	MW	SD	MW	SD
NSB	466	188	466	188	466	188	741	141
1.00E-08							2761	178
3.00E-08							8256	708
1.00E-07							35265	2947
3.00E-07	760	255	491	70	874	475	86859	6139
1.00E-06	2859	455	657	65	589	70	106252	30018
3.00E-06	24498	2290	1028	342	672	88	143232	14064
1.00E-05	61397	7853	3292	556	709	163	150989	24245
3.00E-05	58790	2055	22133	4202	3271	585		
1.00E-04			29600	6936	11322	1668		

5

The angiotensin II receptor antagonist telmisartan brought about a particularly potent activation of the PPARgamma pathway in the PPARgamma reporter cell line. Activation by other angiotensin II receptor antagonists such as losartan and irbesartan took place only at higher test concentrations and to a lesser extent.

10

Example 4: **Examples of formulations**

Tablet 1

15

Tablets having the following composition were obtained by direct compression of the telmisartan sodium salt with excipients and magnesium stearate:

5	Ingredients:	mg
	telmisartan sodium salt	41.708
	mannitol	49.542
	microcrystalline cellulose	50.000
	croscarmellose sodium salt	5.000
10	magnesium stearate	3.750
	total	250.000

Tablet 2

Tablets having the following composition were obtained by direct compression of
15 the telmisartan sodium salt with excipients and magnesium stearate:

	Ingredients:	mg
	telmisartan sodium salt	83.417
	sorbitol	384.083
20	polyvidone K25	25.000
	magnesium stearate	7.500
	total	500.000

Tablet 3

25 Hydrochlorothiazide, telmisartan sodium salt, sorbitol and red iron oxide were mixed in a free fall blender, passed through a 0.8 mm screen and, after the addition of magnesium stearate, processed in a free fall blender to obtain a powdered mixture.

30 This combination of active substances and excipients was then compressed with a suitable tablet press (e.g. Korsch EK0 or Fette P1200) to form tablets. Tablets with the following composition were obtained, the quantity of telmisartan sodium salt contained in each tablet corresponding to a quantity of 80 mg of the free acid of telmisartan.

Ingredient	mg/tablet	%
telmisartan sodium salt	83.417	13.903
hydrochlorothiazide	12.500	2.083
sorbitol	494.483	82.414
red iron oxide	0.600	0.100
magnesium stearate	9.000	1.500
total	600.000	100.000

5

The telmisartan sodium salt of the tablets of the three batches dissolved in 900 ml of 0.1 M phosphate buffer, pH 7.5, at a rate of $92 \pm 1.5 \%$, $96 \pm 1.8 \%$ and $100 \pm 1.0 \%$, respectively, after 30 minutes' stirring (75 rpm). The hydrochlorothiazide dissolved in 900 ml of 0.1 M HCl (100 rpm) after 30 minutes at a rate of $69 \pm 6.3 \%$, $72 \pm 2.1 \%$ and $78 \pm 1.8 \%$, respectively.

10

Example 5: Preparation of a crystalline telmisartan sodium salt, starting from telmisartan

15

The starting material for preparing crystalline telmisartan sodium salt may be the free acid of telmisartan, which may be obtained by conventional methods (e.g. according to EP-0 502 314).

154.4 g of telmisartan were placed in 308.8 ml of toluene in a suitable reaction vessel, the suspension was combined with 27.8 g of a 44.68% sodium hydroxide solution and 84.9 ml of ethanol and heated to 78 °C for about 30 minutes. Then the mixture was filtered. If large amounts of solid were left in the filter, the filter was, optionally, washed with a mixture of 61.8 ml of toluene and 15.3 ml of ethanol.

25

463.2 ml of toluene were placed in another reaction vessel and refluxed. The filtrate obtained according to the process described above was slowly added at the boiling temperature and simultaneously distilled azeotropically. After it was all

5 added, any solution obtained by washing the filter was also added and again
azeotropic distillation was carried out. The mixture was distilled until a
temperature of 103 °C was obtained. Then the suspension was cooled to ambient
temperature. The crystals were suction filtered, washed with 154.4 ml of toluene
and dried at 60 °C in a circulating air dryer.

10

Yield: 154.6 g (96 %)

Colourless crystals

$C_{33}H_{29}N_4O_2Na \times 0.5H_2O$	calc.: C 72.51	H 5.72	N 10.25
	found: C 72.57	H 5.69	N 10.21

15

**Example 6: Preparation of crystalline telmisartan sodium salt from
telmisartan hydrochloride**

20 *Preparation of telmisartan-hydrochloride:*

411 g of tert.-butyl-4'-[[2-n-propyl-4-methyl-6-(1-methylbenzimidazol-2-yl)-
benzimidazol-1-yl]-methyl]-biphenyl-2-carboxylate were suspended in 822 ml
glacial acetic acid and combined with 213 g concentrated aqueous hydrochloric
acid (37 %). The mixture was refluxed. About 640 ml of the solvent were distilled
25 off. The residue remaining was slowly combined with about 620 ml of water at 50-
60 °C. This mixture was combined with 20 g activated charcoal (e.g. Norit SX 2
Ultra). The mixture obtained was stirred for about 10 minutes at constant
temperature. After filtering, the residue was washed 3 times with 25 ml of glacial
acetic acid and about 620 ml of water. The filtrate obtained was again heated to
30 about 50-60 °C and combined with about 2 litres of water. After about 12 hours'
stirring at about 23 °C the crystals formed were suction filtered and washed twice
with about 500 ml of water and once with about 900 ml acetone and then dried at
about 60 °C.

35 Yield: 367 g (92.5 %)

- 5 Colourless crystals
melting point: 278 °C

Preparation of crystalline telmisartan sodium salt from telmisartan-hydrochloride:
55.1 g telmisartan-hydrochloride were taken up in 110.2 ml of toluene, 5.5 ml of
10 water and 55.1 ml isopropanol. This mixture was combined with 36.9 g sodium
methoxide (30 % in methanol) and 2.75 g activated charcoal (e.g. Norit SX 2
Ultra). Then, the mixture was heated to about 75 °C. About 50 ml of the solvent
mixture were distilled off at constant temperature within about 30 minutes. The
suspension obtained was filtered. The residue was washed with about 20 ml of
15 toluene. The filtrate was combined with about 5 ml of water and about 150 ml of
toluene. The mixture obtained was refluxed. About 150 ml of solvent mixture
were distilled off azeotropically (up to 102 °C). The mixture was left to crystallise
for 1 hour at 100 °C. The crystals were suction filtered, washed with about 50 ml
of toluene and dried at about 60 °C.

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Yield: 53.6 g (99 %)

Colourless crystals

$C_{33}H_{29}N_4O_2Na \cdot 0.5H_2O$	calc.:	C 72.51	H 5.72	N 10.25
	found:	C 72.44	H 5.68	N 10.20

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The present invention is not to be limited in scope by the specific embodiments
described herein, which are intended as single illustrations of individual aspects of
the invention, and functionally equivalent methods and components are within the
scope of the invention. Indeed, various modifications of the invention, in addition
30 to those shown and described herein will become apparent to those skilled in the
art from the foregoing description and accompanying drawings. Such
modifications are intended to fall within the scope of the appended claims.
Various patent applications and publications are cited herein, the disclosures of
which are incorporated by reference in their entireties.